WEST Search History

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DATE: Friday, January 09, 2004

Hide?	Set Name	Query	Hit Count
	DB=USPT; THES=ASSIGNEE; PLUR=YES; OP=ADJ		
☐	L5	L4 and VEGF	165
☐	L4	L3 and chimeric adj antibody	165
☐	L3	L2	933
	DB=PGPB,USPT,EPAB,JPAB,DWPI; THES=ASSIGNEE; PLUR=YES; OP=ADJ		
☐	L2	L1 and angiogenesis	3801
☐	L1	VEGF and antibody	5277

END OF SEARCH HISTORY

h e b b cg b chh e g f c e b

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=> scFVs
L1          291 SCFVS

=> L1 and VEGF
           8662 VEGF
             95 VEGFS
           8666 VEGF
                  (VEGF OR VEGFS)
L2            3 L1 AND VEGF

=> L1 and gene therapy
         827526 GENE
         311930 GENES
         875641 GENE
                  (GENE OR GENES)
         212440 THERAPY
          14097 THERAPIES
         220053 THERAPY
                  (THERAPY OR THERAPIES)
          32409 GENE THERAPY
                  (GENE(W)THERAPY)
L3           20 L1 AND GENE THERAPY

=> DIS L3 1- IBIB IABS
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NEWS	4	DEC 08	INPADOC: Legal Status data reloaded
NEWS	5	SEP 29	DISSABS now available on STN
NEWS	6	OCT 10	PCTFULL: Two new display fields added
NEWS	7	OCT 21	BIOSIS file reloaded and enhanced
NEWS	8	OCT 28	BIOSIS file segment of TOXCENTER reloaded and enhanced
NEWS	9	NOV 24	MSDS-CCOHS file reloaded
NEWS	10	DEC 08	CABA reloaded with left truncation
NEWS	11	DEC 08	IMS file names changed
NEWS	12	DEC 09	Experimental property data collected by CAS now available in REGISTRY
NEWS	13	DEC 09	STN Entry Date available for display in REGISTRY and CA/CAplus
NEWS	14	DEC 17	DGENE: Two new display fields added
NEWS	15	DEC 18	BIOTECHNO no longer updated
NEWS	16	DEC 19	CROPU no longer updated; subscriber discount no longer available
NEWS	17	DEC 22	Additional INPI reactions and pre-1907 documents added to CAS databases
NEWS	18	DEC 22	IFIPAT/IFIUDB/IFICDB reloaded with new data and search fields
NEWS	19	DEC 22	ABI-INFORM now available on STN

NEWS EXPRESS	DECEMBER 28 CURRENT WINDOWS VERSION IS V7.00, CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP), AND CURRENT DISCOVER FILE IS DATED 23 SEPTEMBER 2003
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FILE COVERS 1907 - 9 Jan 2004  VOL 140 ISS 3
FILE LAST UPDATED: 8 Jan 2004  (20040108/ED)

  This file contains CAS Registry Numbers for easy and accurate
  substance identification.


=> VEGF
          8655 VEGF
            95 VEGFS
L1        8659 VEGF
                 (VEGF OR VEGFS)

=> chimeric (w) antibody
         37684 CHIMERIC
            24 CHIMERICS
         37693 CHIMERIC
                 (CHIMERIC OR CHIMERICS)
        256291 ANTIBODY
        283989 ANTIBODIES
        390476 ANTIBODY
                 (ANTIBODY OR ANTIBODIES)
L2        971 CHIMERIC (W) ANTIBODY

=> L1 and L2
L3         14 L1 AND L2

=> treatment and L1
       1862991 TREATMENT
        171435 TREATMENTS
       1955694 TREATMENT
                 (TREATMENT OR TREATMENTS)
L4        2003 TREATMENT AND L1

=> L4 and L2
L5            3 L4 AND L2

=> DIS L5 1- IBIB IABS
YOU HAVE REQUESTED DATA FROM 3 ANSWERS - CONTINUE? Y/(N):Y
THE ESTIMATED COST FOR THIS REQUEST IS 7.62 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:Y

L5   ANSWER 1 OF 3  CAPLUS  COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:         2003:335256  CAPLUS
DOCUMENT NUMBER:          138:352765
```

TITLE: Antibody or immunoadhesin having Fc region for diagnosis and **treatment** of cancer, autoimmune disease, inflammation, or infection
INVENTOR(S): Presta, Leonard G.
PATENT ASSIGNEE(S): Genentech, Inc., USA
SOURCE: PCT Int. Appl., 139 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003035835	A2	20030501	WO 2002-US33739	20021022
WO 2003035835	A3	20031016		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003157108	A1	20030821	US 2002-277370	20021022

PRIORITY APPLN. INFO.: US 2001-337642P P 20011025
US 2002-347694P P 20020109

ABSTRACT:
The present invention concerns compns. comprising a glycoprotein having an Fc region, wherein about 80-100% of the glycoprotein in the compn. comprises a mature core carbohydrate structure which lacks fucose, attached to the Fc region of the glycoprotein. The preferred glycoprotein is an antibody or immunoadhesin. The antibody or immunoadhesin is a chimeric, humanized or human antibody or immunoadhesin. The antibody or immunoadhesin is specific to B cell surface marker, ErbB receptor, tumor antigen or angiogenic factor, such as CD20, HER2, **VEGF**, CD40 or prostate stem cell antigen. The antibody or immunoadhesin is useful for treating cancer, autoimmune disease, inflammatory disease, infection, or condition where removal of cells or tissue is desired.

L5 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2001:338563 CAPLUS
DOCUMENT NUMBER: 134:348629
TITLE: Modulation of eNOS activity using **VEGF**, a variant, or **VEGF** receptor agonists and therapeutic uses thereof
INVENTOR(S): Shen, Ben-Quan; Zioncheck, Thomas
PATENT ASSIGNEE(S): Genentech, Inc., USA
SOURCE: PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001032695	A2	20010510	WO 2000-US30294	20001102
WO 2001032695	A3	20020214		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,

CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

EP 1225910 A2 20020731 EP 2000-980281 20001102
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

JP 2003513105 T2 20030408 JP 2001-535394 20001102

PRIORITY APPLN. INFO.: US 1999-163132P P 19991102
WO 2000-US30294 W 20001102

ABSTRACT:
The present invention provides uses of **VEGF**, a variant, or ***VEGF*** receptor agonists for the up-regulation of eNOS expression and activity. **VEGF**, its variants, and **VEGF** receptor agonists are useful in the **treatment** of or prevention from hypertension, diabetes, angina, thrombosis, atherosclerosis, heart failure, and other conditions or disorders wherein nitric oxide is an important regulator. Methods of prepg. the variants are also disclosed in the patent.

L5 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2000:911535 CAPLUS
DOCUMENT NUMBER: 134:85128
TITLE: Diagnostics and remedies for diseases with participation of macrocytes/macrophages
INVENTOR(S): Shitara, Kenya; Shibuya, Masabumi
PATENT ASSIGNEE(S): Kyowa Hakko Kogyo Co., Ltd., Japan
SOURCE: PCT Int. Appl., 163 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000079275	A1	20001228	WO 2000-JP3957	20000616

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR,
CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU,
ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV,
MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE,
SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA,
ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

EP 1199565 A1 20020424 EP 2000-937283 20000616
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI, LT, LV, FI, RO, MK, CY, AL

PRIORITY APPLN. INFO.: JP 1999-171709 A 19990617
WO 2000-JP3957 W 20000616

ABSTRACT:
Diagnostics and remedies for inflammatory diseases, delayed hypersensitivity, malignant tumor and arteriosclerosis which contain, as the active ingredient, a substance binding to human **VEGF** receptor Flt-1 or a substance inhibiting signal transduction mediated by human **VEGF** receptor Flt-1. The human **VEGF** receptor Flt-1-binding substance is a monoclonal or polyclonal antibody, **chimeric antibody**, or antibody fragment.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> DIS L3 1- IBIB IABS
YOU HAVE REQUESTED DATA FROM 14 ANSWERS - CONTINUE? Y/(N):Y
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L3 ANSWER 1 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:972195 CAPLUS
DOCUMENT NUMBER: 140:26922
TITLE: Chimeric/humanized antibodies comprising viral coat protein, peptide tag and linkers for screening target antigen-binding polypeptides as therapeutics and reagents
INVENTOR(S): Fuh, Germaine G.; Sidhu, Sachdev S.
PATENT ASSIGNEE(S): Genentech, Inc., USA
SOURCE: PCT Int. Appl., 205 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003102157	A2	20031211	WO 2003-US17545	20030603

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 2002-385338P P 20020603
US 2003-463656P P 20030416

ABSTRACT:
The invention provides comprising variant amino acids in CDRs of antibody variable domains. These polypeptides provide a source of great sequence diversity that can be used as a source for identifying novel antigen binding polypeptides. The target antigen is **VEGF**, IGF-1 or Her-2. The invention also provides these polypeptides as fusion polypeptides to heterologous polypeptides such as at least a portion of phage or viral coat proteins, tags, and linkers. The viral coat protein consists of protein pIII, major coat protein pVIII, Soc, Hoc, gpD, pv1 or variant; and the peptide tag is gD, c-myc, poly-His, fluorescence protein, or .beta. galactosidase. Libraries comprising a plurality of these polypeptides are also provided. In addn., methods of and compns. for generating and using these polypeptides and libraries are provided.

L3 ANSWER 2 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:863449 CAPLUS
DOCUMENT NUMBER: 139:336930
TITLE: Antibodies specific to KDR/Flk-1 phosphorylated at tyrosine 1214, and its uses in drug screening and therapy
INVENTOR(S): Shibuya, Masashi; Furuya, Akiko; Shitara, Kenya
PATENT ASSIGNEE(S): Kyowa Hakko Kogyo Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 23 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2003310276	A2	20031105	JP 2002-129072	20020430

PRIORITY APPLN. INFO.: JP 2002-129072 20020430

ABSTRACT:
Antibodies specific to vascular endothelial growth factor receptor KDR/Flk-1 phosphorylated at tyrosine at position 1214 (Y1214), and use in various therapeutic and drug screening applications, are disclosed. Various angiogenesis-related methods using this substance are provided: a method for inhibiting the signal transduction of KDR/Flk-1; a method for inhibiting cell proliferation; a method for inhibiting angiogenesis; a method for screening a cell proliferation inhibitor; a method for screening an angiogenesis inhibitor; a method for screening a substance inhibiting the signal transduction of KDR/Flk-1; a method for judging whether or not a test substance inhibits the signal transduction of KDR/Flk-1; a method for screening a substance inhibiting the phosphorylation at tyrosine at the 1214-position of KDR/Flk-1.

L3 ANSWER 3 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:656789 CAPLUS
DOCUMENT NUMBER: 139:196277
TITLE: Antibody variants with faster antigen association rates for diagnostics and therapeutic uses
INVENTOR(S): Lowman, Henry B.; Marvin, Jonathan S.
PATENT ASSIGNEE(S): Genentech, Inc., USA
SOURCE: PCT Int. Appl., 81 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003068801	A2	20030821	WO 2003-US4184	20030211
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
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US 2003224397	A1	20031204	US 2003-364953	20030211

PRIORITY APPLN. INFO.: US 2002-355895P P 20020211
US 2002-409685P P 20020910

ABSTRACT:
Antibody variants with higher affinity to antigen are disclosed. The antibody variants have one or more amino acid alteration(s) in or adjacent to at least one hypervariable region thereof which increase charge complementarity between the antibody variant and an antigen to which it binds. Variants of anti-***VEGF*** antibody Y0101, anti-tissue factor antibody D3H44 and anti-HER2 antibody 4D5 were prepd. and tested.

L3 ANSWER 4 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:335256 CAPLUS
DOCUMENT NUMBER: 138:352765
TITLE: Antibody or immunoadhesin having Fc region for diagnosis and treatment of cancer, autoimmune disease, inflammation, or infection
INVENTOR(S): Presta, Leonard G.
PATENT ASSIGNEE(S): Genentech, Inc., USA
SOURCE: PCT Int. Appl., 139 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003035835	A2	20030501	WO 2002-US33739	20021022
WO 2003035835	A3	20031016		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003157108	A1	20030821	US 2002-277370	20021022

PRIORITY APPLN. INFO.: US 2001-337642P P 20011025
US 2002-347694P P 20020109

ABSTRACT:
The present invention concerns compns. comprising a glycoprotein having an Fc region, wherein about 80-100% of the glycoprotein in the compn. comprises a mature core carbohydrate structure which lacks fucose, attached to the Fc region of the glycoprotein. The preferred glycoprotein is an antibody or immunoadhesin. The antibody or immunoadhesin is a chimeric, humanized or human antibody or immunoadhesin. The antibody or immunoadhesin is specific to B cell surface marker, ErbB receptor, tumor antigen or angiogenic factor, such as CD20, HER2, **VEGF**, CD40 or prostate stem cell antigen. The antibody or immunoadhesin is useful for treating cancer, autoimmune disease, inflammatory disease, infection, or condition where removal of cells or tissue is desired.

L3 ANSWER 5 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:273850 CAPLUS
DOCUMENT NUMBER: 139:336677
TITLE: A Phase I Study of Anti-Kinase Insert Domain-containing Receptor Antibody, IMC-1C11, in Patients with Liver Metastases from Colorectal Carcinoma
AUTHOR(S): Posey, James A.; Ng, Thian C.; Yang, Baolian; Khazaeli, M. B.; Carpenter, Mark D.; Fox, Floyd; Needle, Mike; Waksal, Harlan; LoBuglio, Albert F.
CORPORATE SOURCE: Comprehensive Cancer Center, Division of Hematology/Oncology, Departments of Medicine and Radiation Oncology, University of Alabama at Birmingham, Birmingham, AL, 35294-3300, USA
SOURCE: Clinical Cancer Research (2003), 9(4), 1323-1332
CODEN: CCREF4; ISSN: 1078-0432

PUBLISHER: American Association for Cancer Research
DOCUMENT TYPE: Journal
LANGUAGE: English
ABSTRACT:
Angiogenesis plays an important role in colorectal cancer progression. Stimulation of vascular endothelial growth factor receptor (VEGFR), a transmembrane glycoprotein, results in endothelial mitogenesis. Within this family of receptors, VEGFR2/kinase-insert-domain-contg. receptor (KDR) appear to be principally up-regulated during tumorigenesis. A chimeric anti-KDR antibody, IMC-1C11, blocks VEGFR-KDR interaction and inhibits VEGFR-induced endothelial cell proliferation. This trial seeks to assess the safety, tolerability and feasibility of targeting an important pathway in tumorigenesis. In a dose-escalation, single-agent study of IMC-1C11, we enrolled 14 patients with colorectal carcinoma and hepatic metastases. Safety-, pharmacokinetic-, immunogenicity-, and magnetic resonance imaging-assessed alteration of vascular effects of IMC-1C11 were evaluated in this trial. IMC-1C11 was infused weekly at 0.2 mg/kg (n = 3), 0.6 mg/kg (n = 4), 2.0 mg/kg (n = 3), and 4.0 mg/kg (n = 4) for 4 wk, which constituted a cycle. No grade-3 or -4 IMC-1C11-related toxicities were obsd. Minor grade-1 bleeding events were obsd. in four patients [0.2 mg/kg (n = 1) and 0.6 mg/kg (n = 3)]. Each resolved quickly and required no intervention. The starting dose of IMC-1C11 was selected to achieve a Cmax of .apprx.5 .mu.g/mL. This concn. prevented KDR phosphorylation in vitro. Pharmacokinetic anal. demonstrated that the plasma t1/2 and Cmax were dose dependent with a plasma t1/2 of 67 .+-. 3 h at the 4-mg/kg dose level. Human anti-**chimeric** ***antibodies*** were detected in 7 of 14 patients. The antibodies to IMC-1C11 inhibited the circulation of the agent in two patients. One patient had prolonged stable disease for seven cycles (28 wk). The mean changes in tumor-influx vol.-transfer const. kin (min-1) and enhancement factor after 4 wk of therapy were significantly decreased compared with pretreatment values in 11 patients. IMC-1C11 was both safe and well tolerated. Drug levels of IMC-1C11 were reliably predicted. Further clin. investigation of anti-VEGFR/KDR agents is warranted.

REFERENCE COUNT: 47 THERE ARE 47 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 6 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2002:814159 CAPLUS
DOCUMENT NUMBER: 137:336728
TITLE: **Chimeric antibodies** and fragments or variants specific to vascular endothelial growth factor 2 for diagnosing, prognosing and treating infection, inflammation, cancer and autoimmune disease
INVENTOR(S): Rosen, Craig A.; Albert, Vivian R.; Ruben, Steven M.; Wager, Ruth E.
PATENT ASSIGNEE(S): Human Genome Sciences, Inc., USA
SOURCE: PCT Int. Appl., 407 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002083704	A1	20021024	WO 2002-US11474	20020412

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003175274 A1 20030918 US 2002-120414 20020412

WO 2003097660 A1 20031127 WO 2002-US26246 20020819

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 2001-283385P P 20010413
US 2002-350366P P 20020124
WO 2002-US11474 A 20020412

ABSTRACT:
Disclosed are anti-**VEGF**-2 agonistic or antagonistic human or humanized antibodies, antibody fragments, or variants thereof. Also provided are processes for producing such antibodies. The present invention relates to methods and compns. for preventing, treating or ameliorating a disease or disorder comprising administering to an animal, preferably a human, an effective amt. of one or more **VEGF**-2 antibodies or fragments or variants thereof. The disease is selected from inflammatory diseases, proliferative diseases, tumors, metastasis, breast cancer, brain cancer, prostate cancer, colon cancer, lymphangioma, infections, Kaposi's sarcoma, autoimmune diseases, rheumatoid arthritis, psoriasis, diabetic retinopathy, and other diseases assocd. with aberrant or lack of **VEGF**-2 expression.

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 7 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2002:709689 CAPLUS
DOCUMENT NUMBER: 137:211928
TITLE: Construction of eukaryotic expression system and its uses for antibody cloning
INVENTOR(S): Yang, Zhihua; Ran, Yuliang
PATENT ASSIGNEE(S): Inst. of Tumors, Tumor Hospital, Chinese Academy of Medical Sciences, Peop. Rep. China
SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu, 31 pp. CODEN: CNXXEV
DOCUMENT TYPE: Patent
LANGUAGE: Chinese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 1328157	A	20011226	CN 2000-108065	20000609

PRIORITY APPLN. INFO.: CN 2000-108065 20000609

ABSTRACT:
The invention relates to construction of plasmid vector for expression of antibody in eukaryote. The vector contains selective marker gene (such as aminoglycoside phosphate transferase, thymidine kinase, hygromycin B phosphate transferase, xanthine-guanine phosphoribosyltransferase, or asparagine synthase) and extensible selective marker gene (such as dihydrofolic acid reductase (dhfr) or glutamine synthase). The vector also contains weak promotor (such as 72 bp fragment-deleted SV40) for driving the expression of the said marker genes. The vector also contains strong promotor (such as PhCMV-IE, PSV40-E, or PRSV-LTR) for driving the expression of antibody gene.

The vector also contains enhancer sequence (such as 5'-non-translational region SP163 of mouse vascular endothelial growth factor). The vector further contains strong translation terminator (such as BGH polyA or SV40 polyA). The eukaryotic expression system is constructed by successively prepg. general cloning vectors (such as pYR-GCVH and pYR-GCVL) of variable region of antibody, intermediate expression vectors (such as pYR-SV2-rdhfr and pYR-SV2-rneo) of antibody, and general eukaryotic expression vector (such as pYR-GSEVH, pYR-GSEVL, pYR-GCEVH, and pYR-GCEVL). The eukaryotic expression system is used to prepn. and prodn. of antibodies (such as **chimeric antibody** , modified antibody, humanized antibody, small mol. antibody, intracellular antibody, double-specific antibody, and other derivs.). The **chimeric** ***antibody*** of human **VEGF** and small mol. antibody of human carcino-embryonic antigen (CEA) were prepd. by using the eukaryotic expression system in CHO-dhfr- cell.

L3 ANSWER 8 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2002:695813 CAPLUS
DOCUMENT NUMBER: 137:231369
TITLE: Anti-VEGFR antibodies in combination with anti-EGFR antibodies, chemotherapeutic agent or radiotherapeutic agent for inhibiting tumor growth
PATENT ASSIGNEE(S): Imclone Systems Incorporated, USA; Rockwell, Patricia; Goldstein, Neil I.
SOURCE: PCT Int. Appl., 151 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 6
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002070008	A1	20020912	WO 2002-US6762	20020304
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2003103973	A1	20030605	US 2001-798689	20010302

PRIORITY APPLN. INFO.:

US 2001-798689	A	20010302
US 1994-196041	B2	19940210
US 1994-326552	A1	19941020
US 1995-476533	B2	19950607
US 1996-706804	A2	19960903
US 1997-967113	A1	19971110
US 1999-401163	A2	19990922

ABSTRACT:
The present invention provides a method of reducing or inhibiting tumor growth in a mammal comprising treating the mammal with an effective amt. of a combination of a **VEGF** receptor antagonist and radiation, chemotherapy, and/or an addnl. receptor antagonist (e.g. epidermal growth factor receptor antagonist). The **VEGF** receptor antagonists are anti-VEGFR antibodies, fragments, humanized or **chimeric** ***antibodies*** .

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 9 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2002:409126 CAPLUS
DOCUMENT NUMBER: 137:5011
TITLE: Antibodies specific to human KDR **VEGF** receptor for inhibiting angiogenesis and tumor growth
INVENTOR(S): Zhu, Zhenping; Witte, Larry
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 34 pp., Cont. of U.S. Ser. No. 493,539.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002064528	A1	20020530	US 2001-976787	20011012

PRIORITY APPLN. INFO.: US 2000-493539 A1 20000128
ABSTRACT:
The invention provides an Ig. mol. which binds KDR with an affinity comparable to human **VEGF**, and that neutralizes activation of KDR. Ig. mols. include monovalent single chain antibodies, multivalent single chain antibodies, diabodies, triabodies, antibodies, humanized antibodies and ***chimeric*** **antibodies**. The invention further provides nucleic acid mols. that encode these Ig. mols. The invention also provides a method of making the Ig. mols. mentioned above. The invention further provides a method of neutralizing the activation of KDR, a method of inhibiting angiogenesis in a mammal and a method of inhibiting tumor growth in a mammal with such Ig. mols.

L3 ANSWER 10 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2001:338563 CAPLUS
DOCUMENT NUMBER: 134:348629
TITLE: Modulation of eNOS activity using **VEGF**, a variant, or **VEGF** receptor agonists and therapeutic uses thereof
INVENTOR(S): Shen, Ben-Quan; Zioncheck, Thomas
PATENT ASSIGNEE(S): Genentech, Inc., USA
SOURCE: PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001032695	A2	20010510	WO 2000-US30294	20001102
WO 2001032695	A3	20020214		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
EP 1225910	A2	20020731	EP 2000-980281	20001102
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
JP 2003513105	T2	20030408	JP 2001-535394	20001102

PRIORITY APPLN. INFO.: US 1999-163132P P 19991102
WO 2000-US30294 W 20001102

ABSTRACT:
The present invention provides uses of **VEGF**, a variant, or ***VEGF*** receptor agonists for the up-regulation of eNOS expression and activity. **VEGF**, its variants, and **VEGF** receptor agonists are useful in the treatment of or prevention from hypertension, diabetes, angina, thrombosis, atherosclerosis, heart failure, and other conditions or disorders wherein nitric oxide is an important regulator. Methods of prepg. the variants are also disclosed in the patent.

L3 ANSWER 11 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2000:911535 CAPLUS
DOCUMENT NUMBER: 134:85128
TITLE: Diagnostics and remedies for diseases with participation of macrocytes/macrophages
INVENTOR(S): Shitara, Kenya; Shibuya, Masabumi
PATENT ASSIGNEE(S): Kyowa Hakko Kogyo Co., Ltd., Japan
SOURCE: PCT Int. Appl., 163 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000079275	A1	20001228	WO 2000-JP3957	20000616
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
EP 1199565	A1	20020424	EP 2000-937283	20000616
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL				

PRIORITY APPLN. INFO.: JP 1999-171709 A 19990617
WO 2000-JP3957 W 20000616

ABSTRACT:
Diagnostics and remedies for inflammatory diseases, delayed hypersensitivity, malignant tumor and arteriosclerosis which contain, as the active ingredient, a substance binding to human **VEGF** receptor Flt-1 or a substance inhibiting signal transduction mediated by human **VEGF** receptor Flt-1. The human **VEGF** receptor Flt-1-binding substance is a monoclonal or polyclonal antibody, **chimeric antibody**, or antibody fragment.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 12 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2000:219697 CAPLUS
DOCUMENT NUMBER: 133:118695
TITLE: Construction and expression of a chimeric murine-human antibody against VEGF165
AUTHOR(S): Guo, Wenzhong; Yang, Zhihua; Ran, Yuliang; Wang, Guiqi; Liu, Jun; Sun, Lixui; Dong, Zhiwei
CORPORATE SOURCE: Cancer Inst., Chinese Acad. Med. Sci., Beijing, 100021, Peop. Rep. China
SOURCE: Zhonghua Weishengwuxue He Mianyixue Zazhi (2000), 20(1), 45-48

CODEN: ZWMZDP; ISSN: 0254-5101
PUBLISHER: Weishenbu Beijing Shengwu Zhipin Yanjiuso
DOCUMENT TYPE: Journal
LANGUAGE: Chinese
ABSTRACT:
Objective To construct a chimeric murine-human antibody against VEGF165. Methods The murine genes encoding the variable regions of light and heavy chains of monoclonal antibody VmD11 against VEGF165 were cloned into expressional vectors of pAcyc-neo-C.kappa. and psv2-gpt-C.gamma.1 sep. The resulted vectors were transferred into murine-myeloma-cells to express ***chimeric*** **antibodies**. The **chimeric antibody** in the cultural supernatant of transfected cells was detected by indirect ELISA and its humanized character and specificity against VEGF165 was confirmed by Western blot, RT-PCR and competitive ELISA. Results The **chimeric** ***antibody*** against human VEGF165 was detected in the cultural supernatant of transfected myeloma cells with a yield of 10.mu.g/L. Specificity of the ***chimeric*** **antibody** was proved by Western blot and competitive ELISA, the expression of chimeric gene was confirmed by RT-PRC. Conclusion Chimeric mouse-human antibody against VEGF165 was successfully expressed in myeloma cells of mouse.

L3 ANSWER 13 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2000:7095 CAPLUS
DOCUMENT NUMBER: 132:320697
TITLE: Construction of anti-human VEGF165 **chimeric antibodies** and expression in eukaryotic cells
AUTHOR(S): Ran, Yuliang; Yang, Zhihua; Wang, Guiqi; Liu, Jun; Sun, Lixin; Dong, Zhiwei
CORPORATE SOURCE: Cancer Institute (Hospital), Chinese Academy of Medical Sciences, Peking Union Medical College, Beijing, 100021, Peop. Rep. China
SOURCE: Zhonghua Zhongliu Zazhi (1999), 21(6), 412-415
CODEN: CCLCDY; ISSN: 0253-3766
PUBLISHER: Zhongguo Yixue Kexueyuan Zhongliu Yanjiuso, Zhongliu Yiyuan
DOCUMENT TYPE: Journal
LANGUAGE: Chinese
ABSTRACT:
The chimeric anti-vascular endothelial growth factor 165 antibody gene was expressed in eukaryotic cells. The variable region genes of light and heavy chains from mouse anti-human VEGF165 monoclonal antibody VmD11 were cloned into eukaryotic expression vectors and transfected into dihydrofolated reductase-deficient Chinese hamster ovary (CHO-dhfr-) cells to express ***chimeric*** **antibody**. The antibody expressed was examd. for the presence of human const. regions and specificity against human VEGF165. The ***chimeric*** **antibody** with human const. regions and specificity against human **VEGF** was detected in the culture supernatant of transfected CHO cells by ELISA and Western blot. The **chimeric** ***antibody*** gene was also detected at mRNA level by RT-PCR. The mouse anti-human VEGF165 monoclonal antibody is successfully humanized and expressed in eukaryotic cells.

L3 ANSWER 14 OF 14 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 1995:826808 CAPLUS
DOCUMENT NUMBER: 123:225950
TITLE: Monoclonal antibodies specific to **VEGF** receptors and uses thereof
INVENTOR(S): Rockwell, Patricia; Goldstein, Neil I.
PATENT ASSIGNEE(S): Imclone Systems Incorp., USA
SOURCE: PCT Int. Appl., 67 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent

LANGUAGE: English
FAMILY ACC. NUM. COUNT: 6
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9521868	A1	19950817	WO 1995-US1678	19950210
W: AU, CA, CN, FI, HU, JP, KR, NO, RU				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
US 5840301	A	19981124	US 1994-326552	19941020
AU 9519147	A1	19950829	AU 1995-19147	19950210
EP 741748	A1	19961113	EP 1995-911659	19950210
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				

PRIORITY APPLN. INFO.: US 1994-196041 A 19940210
US 1994-326552 A 19941020
WO 1995-US1678 W 19950210

ABSTRACT:
Monoclonal antibodies that specifically bind to an extracellular domain mammalian or human vascular endothelial growth factor (**VEGF**) receptor and neutralize activation of the receptor in endothelial or tumor cells are provided. The monoclonal antibodies are used in combination with a chemotherapeutic agent, e.g. doxorubicin, cisplatin, or taxol, for inhibition of tumor growth or angiogenesis. **Chimeric antibodies** comprising murine variable region and human const. region, polypeptides contg. the variable region of the monoclonal antibody, and nuclei acid sequences encoding the polypeptide are all claimed.

- Search History will be lost after eight hours of inactivity.
- To combine searches use # before search number, e.g., #2 AND #6.
- Search numbers may not be continuous; all searches are represented.

Search	Most Recent Queries	Time	Result
#23	Search **VEGF flt-1 and chimeric antibody**	14:52:56	9
#15	Search **VEGF flt-1 and chimeric**	14:25:11	22
#14	Search **VEGF flt-1**	14:19:35	863
#10	Search **Rockwell P and VEGF**	14:15:21	9
#7	Search **Wang M and MSP**	13:41:04	24
#6	Search **Wang M**	13:40:55	3085
#5	Search **Wang M-H**	13:39:57	34
#3	Search **Ssalvetti A 1998 and AAV**	10:32:27	108
#1	Search **Zhou X 1998 and AAV**	10:31:04	1

Clear History

- Search History will be lost after eight hours of inactivity.
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- Search numbers may not be continuous; all searches are represented.

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Related Resources

Search	Most Recent Queries	Time	Result
#17	Search **scFVs and gene therapy**	12:25:19	13
#14	Search **scFVs and VEGF**	12:12:10	3
#2	Search **VEGF neutralizing antibody and cancer treatment**	11:57:15	56
#6	Search **Rosenberg SA[au]** Limits: **Clinical Trial**	11:27:00	114
#4	Search **VEGF neutralizing antibody and cancer treatment** Field: **All Fields,** Limits: **Clinical Trial**	11:11:24	3
#1	Search **VEGF neutralizing antibody**	11:09:41	226

Clear History

1: Hybridoma. 1995 Oct;14(5):475-80. Related Articles, Links

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Related Resources

Isolation and characterization of neutralizing monoclonal antibodies to human vascular endothelial growth factor/vascular permeability factor121 (VEGF/VPF121).

Asano M, Yukita A, Matsumoto T, Matsuo K, Kondo S, Suzuki H.

Bioscience Research Department, Tsukuba Research Laboratory, Toagosei Co. Ltd., Ibaraki, Japan.

We have established monoclonal antibodies (MAbs) against human vascular endothelial growth factor/vascular permeability factor121 (VEGF/VPF121). Two (MV101 and MV303) of the 28 MAbs neutralized the mitogenic activity of VEGF/VPF121 on human umbilical vein endothelial cells (HUVEC) in a dose-dependent manner. Both of the MAbs reacted to VEGF/VPF121 and also VEGF/VPF165 with somewhat different binding properties in a sandwich-type enzyme-linked immunosorbent assay (ELSIA). The binding of MV101 and MV303 to VEGF/VPF121 was competitive, but MV415, another anti-VEGF/VPF121 MAb without neutralizing activity, did not complete with either of the antibodies. MV101 and MV303 specifically recognized the native form of VEGF/VPF121 and VEGF/VPF165 in Western blotting. They did not react with VEGF/VPF when the antigens were fractionated under reducing conditions. These observations suggested that MV101 and MV303 might recognize the epitopes closely located on the configuration of VEGF/VPF121 molecule and the epitopes recognized by MV101 and MV303 may play an important role in the VEGF/VPF-receptor signal transduction. These MAbs significantly suppressed the growth of a human hepatoma, PLC/PRF/5, in vivo.

PMID: 8575796 [PubMed - indexed for MEDLINE]

1: J Urol. 1999 Mar;161(3):960-3.

Neutralizing anti-vascular endothelial growth factor antibody inhibits further growth of established prostate cancer and metastases in a pre-clinical model.

Melnyk O, Zimmerman M, Kim KJ, Shuman M.

Cancer Research Institute, University of California at San Francisco, USA.

PURPOSE: The formation of new blood vessels from the pre-existing vasculature is necessary for support of primary tumor growth and appears coincident with the development of metastasis. In previous studies, inhibition of vascular endothelial growth factor (VEGF), a potent angiogenic factor and mediator of vascular permeability, inhibited tumor neovascularization with consequent inhibition of both primary tumor growth and micrometastases when administered at the time of tumor inoculation. In the present study, we examined the effect of inhibiting VEGF on primary tumor growth and metastases in an in vivo model of established metastatic prostate cancer. MATERIALS AND METHODS: The human prostate cancer cell line DU-145 was found to secrete VEGF. DU-145.luciferase, a subclone stably transfected with an expression vector encoding the luciferase gene, injected subcutaneously, consistently formed tumors in C.B.-17 scid/scid mice. After 6 weeks, assay of whole lung lysates showed significant luciferase activity, consistent with the presence of micrometastasis. RESULTS: Twice weekly treatment of the animals with a monoclonal anti-VEGF neutralizing antibody, A4.6.1, not only suppressed primary tumor growth, but inhibited metastatic dissemination to the lung. When treatment was delayed until the primary tumors were well-established, further growth was still inhibited, as was the progression of metastatic disease. CONCLUSION: Inhibition of tumor-secreted VEGF by a neutralizing antibody is sufficient to significantly impair prostate tumor growth and its subsequent metastasis in an in vivo model of established advanced prostate cancer. These data suggest a critical role for VEGF in initiation and maintenance of tumor angiogenesis in prostate cancer. Inhibition of VEGF in patients with VEGF-secreting prostate cancers may prove an effective approach for inhibiting disease progression even after micro-metastatic dissemination has occurred.

PMID: 10022734 [PubMed - indexed for MEDLINE]

1: Hum Gene Ther. 1996 Nov 10;7(17):2157-64. Related Articles, Links

Retroviral display of antibody fragments; interdomain spacing strongly influences vector infectivity.

Ager S, Nilson BH, Morling FJ, Peng KW, Cosset FL, Russell SJ.

Cambridge Centre for Protein Engineering, England.

Five different single-chain antibody fragments (scFv) against human cell-surface antigens were displayed on murine ecotropic retroviral vectors by fusing them to the Moloney SU envelope glycoprotein. The spacing between the scFv and the SU glycoprotein was varied by fusing the scFv to residue +7 or to residue +1 of Moloney SU and by inserting linker sequences of different lengths between the domains. All of the chimeric envelopes were efficiently incorporated into vector particles and could bind to human cells through their displayed antibody fragments, but did not infect them. The spacing between the scFvs and the SU glycoproteins had no significant effect on the efficiency of envelope expression or viral incorporation and did not affect the binding properties of the chimeric envelopes, nor did it influence the efficiency of targeted gene delivery to human cells by scFv-displaying vectors. However, on murine fibroblasts the infectivity of vectors incorporating the chimeric envelopes was strongly influenced by the length of the interdomain spacer. The titers were very low when the single-chain antibodies were fused through a tripeptide linker to SU residue +7 and were greatly enhanced (up to 10(5)-fold) when they were fused to SU residue +1 through a heptapeptide linker. These results point to the importance of steric interactions between the domains of chimeric envelope glycoproteins and may have implications for retroviral vector design for human gene therapy.

PMID: 8934229 [PubMed - indexed for MEDLINE]

1: Nat Med. 1997 Nov;3(11):1222-7. Related Articles, Links

Entrez PubMed

PubMed Services

Related Resources

Halting angiogenesis suppresses carcinoma cell invasion.

Skobe M, Rockwell P, Goldstein N, Vosseler S, Fusenig NE.

Division of Carcinogenesis and Differentiation, German Cancer Research Center (DKFZ), Heidelberg.

The importance of angiogenesis in malignant tumor growth has been interpreted mainly in terms of oxygen and nutrient supply. Here we demonstrate its fundamental role for tumor invasion of malignant human keratinocytes in surface transplants on nude mice. Distinct patterns of angiogenesis and vascular endothelial growth factor receptor-2 (VEGFR-2) expression allowed us to distinguish between benign and malignant cells. Functional inactivation of VEGF-R2 by a blocking antibody disrupted ongoing angiogenesis and prevented invasion of malignant cells, without reducing tumor cell proliferation. The reversion of a malignant into a benign phenotype by halting angiogenesis demonstrates a significant function of vascular endothelium for tumor invasion.

PMID: 9359696 [PubMed - indexed for MEDLINE]

Display Abstract Show: 20 Sort Send to Text

Stem Cells, Vol. 19, No. 1, 24-36, January 2001

Anti-VEGFR-2 scFvs for Cell Isolation. Single-Chain Antibodies Recognizing the Human Vascular Endothelial Growth Factor Receptor-2 (VEGFR-2/flk-1) on the Surface of Primary Endothelial Cells and Preselected CD34$^+$ Cells from Cord Blood

Thomas Böldicke[a], Michael Tesar[b], Carsten Griesel[c], Manfred Rohde[b], Hermann-Josef Gröne[e], Johannes Waltenberger[f], Orit Kollet[g], Tsvee Lapidot[g], Avner Yayon[h], Herbert Weich[d]

[a] German Research Centre for Biotechnology, Department of Applied Genetics;
[b] Division of Microbiology;
[c] Department of Biochemical Engineering;
[d] Department of Gene Regulation and Differentiation, Braunschweig, Germany;
[e] DKFZ, Institute of Cellular and Molecular Pathology, Heidelberg, Germany;
[f] Ulm University Medical Center, Department of Internal Medicine II, Ulm, Germany; Weizmann Institute of Science,
[g] Department of Immunology and
[h] Department of Molecular Cell Biology, Rehovot, Israel

Key Words: Immune V-gene phage display library· Single-chain antibody· Human VEGF receptor 2· FACS analysis· Hematopoitic stem cell

Correspondence: Thomas Böldicke, Ph.D., German Research Centre for Biotechnology, Department of Applied Genetics, Mascheroder Weg 1, D-38124 Braunschweig, Germany. Telephone: 49-(0)531-6181-209; Fax: 49-(0)531-6181-202; e-mail: tbo@gbf.de

Abstract

Five specific single-chain antibodies recognizing the human vascular endothelial growth factor receptor-2 (VEGFR-2/KDR) were selected from a V-gene phage display library constructed from mice immunized with the extracellular domain of VEGFR-2 (Ig-like domain 1-7). All five scFv antibodies (A2, A7, B11, G3, and H1) bound to the purified native antigen in enzyme-linked immunosorbent assay and Dot Blot, and showed no crossreactivity to the human VEGF-receptor 1 (VEGFR-1). The selected antibodies recognize a conformation-dependent epitope of the native receptor and do not recognize